

Search Forms

eRed Folder :

Search

Results[Previous Doc](#)[Next Doc](#)[Go to Doc#](#)

User Searches

Preferences

eRed Folder

Logout



Generate Collection

1: Entry 22 of 30

File: JPAB

Mar 22, 1991

PUB-NO: JP403066613A

DOCUMENT-IDENTIFIER: JP 03066613 A

TITLE: FORMATION OF ULTRAFINE PARTICLE OF SLIGHTLY SOLUBLE DRUG

PUBN-DATE: March 22, 1991

INVENTOR-INFORMATION:

NAME

COUNTRY

SAMEJIMA, MASAYOSHI

NODA, KAZUO

KOBAYASHI, YUKIO

OSAWA, TAKASHI

ASSIGNEE-INFORMATION:

NAME

COUNTRY

TANABE SEIYAKU CO LTD

APPL-NO: JP01204132

APPL-DATE: August 4, 1989

INT-CL (IPC): A61K 9/14; A61J 3/02

ABSTRACT:

PURPOSE: To make a slightly soluble drug into ultrafine particles and to improve and promote absorption of slightly soluble drug in digestive tube by blending a low-molecular saccharide with the slightly soluble drug and grinding.

CONSTITUTION: A mixture of a slightly soluble drug and a saccharide (e.g. glucose) or a sugaralcohol is ground by high-speed stirring or impact grinding and the slightly soluble drug is made into ultrafine particles having ≤ 1 micron. 1 pts.wt. of the main drug is used with 1-50 pts.wt. saccharide. The slightly soluble drug has ≤ 5 mg/ml water solubility at 20°C and has insufficient absorption when made into solid preparation by conventional pharmaceutical manufacturing. The drug is nifedipine, indomethacin, cortisone, etc. The ultrafine particles of the drug is contained as an active ingredient and can be pharmaceutically manufactured.

COPYRIGHT: (C)1991, JPO&Japio

[Previous Doc](#)[Next Doc](#)[Go to Doc#](#)